

1       **WHAT IS CLAIMED IS:**

2           **1.** A method for producing multi-gene recombinant vector constructs,

3       which comprises:

4           (1) a multi-gene assembly vector system comprising an acceptor vector

5       and at least two donor vectors; and

6           (2) a DNA recombination system allowing two or more rounds of gene

7       assembly by sequential DNA delivery into the acceptor vector via DNA

8       swapping between the acceptor vector and different donor vectors; and multiple

9       donor vectors will be rotatively used in different rounds of recombination to

10      allow sequential insertion of genes or DNA fragments into the acceptor vector.

11           **2.** The method of claim 1, wherein said acceptor vector comprises:

12           (1) a site RS for DNA recombination;

13           (2) a site S1 located near said RS, which is a cutting site for a homing

14       endonuclease or a restriction endonuclease, or a site for irreversible specific

15       recombination;

16           (3) a selection marker gene that is different from that contained in said

17       donor vector; and

18           (4) a replicon for replication, including those capable of maintaining

19       large plasmids.

20           **3.** The method of claim 1, wherein a donor vector I comprises:

21           (1) a site RS for DNA recombination, which is the same site RS as on

22       said acceptor vector or can form a specific recombination with the RS on said

23       acceptor vector;

24           (2) a site S1 and another site S2, which are cutting sites for homing

1 endonucleases, or for rare-cutting restriction endonucleases, or sites for  
2 irreversible recombination;  
3 (3) a multi-cloning site MCS;  
4 (4) locations of the sites of RS, S1, S2 and MCS on said donor vector in  
5 relative order of RS-S2-MCS-S1; and  
6 (5) a selection marker gene different from that contained in said acceptor  
7 vector.

8 **4.** The method of claim 1, wherein another donor vector II comprises:  
9 (1) a site RS for DNA recombination, which is the same site RS as on  
10 said acceptor vector or can form a specific recombination with the RS on said  
11 acceptor vector;  
12 (2) a site S1 and another site S2, which are cutting sites for homing  
13 endonucleases, or for rare-cutting restriction endonucleases, or sites for  
14 irreversible recombination;  
15 (3) a multi-cloning site MCS;  
16 (4) locations of the sites of RS, S1, S2 and MCS on said donor vector in  
17 relative order of RS-S1-MCS-S2; and  
18 (5) a selection marker gene different from that contained in said acceptor  
19 vector.

20 **5.** The method according to claim 1, wherein said multi-gene assembly  
21 vector system comprising an acceptor vector and donor vector I and donor vector  
22 II is used to carry out two or more cycles of DNA recombination by alternate use  
23 of said donor vector I and donor vector II together with said acceptor vector to  
24 construct multi-gene vector constructs, which recombination process comprises

1 the steps of:

2 (1) cloning of target single genes or gene groups of interest including  
3 any DNA fragments by conventional molecular cloning techniques into the MCS  
4 of said donor vector I or donor vector II, to make the target gene or genes  
5 inserted between the sites S1 and S2;

6 (2) carrying out the first cycle of DNA recombination to recombine the  
7 first target gene or gene groups into said acceptor vector through: (i) *in vivo* or *in*  
8 *vitro* plasmid recombination with said donor vector I containing target gene and  
9 said acceptor vector, upon double-selection of transformants with the selection  
10 marker genes of the plasmids; (ii) removing the backbone sequence of said donor  
11 vector I between the two S1 sites from the integrative plasmid by digestion with  
12 an endonuclease that cut S1 sites, followed by plasmid circularization by ligation  
13 with T4 DNA ligase, and if necessary, with the aid of a double-stranded  
14 oligonucleotide linker compatible to the S1 cutting ends; and (iii) if S1 is an  
15 irreversible recombination site, performing an *in vivo* or *in vitro* recombination  
16 reaction with corresponding recombinase, upon which the backbone sequence of  
17 said donor vector I between the two S1 sites is removed and the ends of the  
18 acceptor vector bearing the inserted gene or gene group are joined to form a  
19 circular plasmid;

20 (3) carrying out the second cycle of DNA recombination to recombine  
21 the second target gene or gene groups into said acceptor vector through: (i) *in*  
22 *vivo* or *in vitro* plasmid recombination with said donor vector II containing the  
23 second gene or gene group and the acceptor vector plasmid obtained from the  
24 step (2), upon double-selection of transformants with the selection marker genes

1 of the plasmids; (ii) removing the backbone sequence of the donor vector II  
2 between the two S2 sites from the integrative plasmid by digestion with an  
3 endonuclease that cut S2 sites followed by plasmid circularization by ligation  
4 with T4 DNA ligase, and if necessary, with the aid of a double-stranded  
5 oligonucleotide linker compatible to the S2 cutting ends; and (iii) if S2 is an  
6 irreversible recombination site, performing an *in vivo* or *in vitro* recombination  
7 reaction with corresponding recombinase, upon which the backbone sequence of  
8 the donor vector II between the two S2 sites is removed and the ends of the  
9 acceptor vector bearing the inserted genes or gene groups are joined to form a  
10 circular plasmid;

11 (4) repeating said step (2) and step (3) with alternate donor vector  
12 plasmids containing target gene or gene group and the acceptor vector plasmid  
13 obtained in the former step, until all target genes or DNA fragments being linked  
14 into the acceptor vector to finish a designed vector construct.

15 **6.** An acceptor vector plasmid according to claim 1, which comprises all  
16 or part of the components shown in Fig. 1A or all or part of the DNA sequence  
17 SEQ ID NO: 1.

18 **7.** A donor vector I plasmid according to claim 1, which comprises all or  
19 part of the components shown in Fig. 1B or all or part of the DNA sequence SEQ  
20 ID NO: 2.

21 **8.** A donor vector II plasmid according to claim 1, which comprises all  
22 or part of the components shown in Fig. 1C or all or part of the DNA sequence  
23 SEQ ID NO: 3.

24 **9.** The application of the method of claim 1, wherein multiple genes or

1 DNA fragments of interest are combined to a vector to create a desired vector  
2 construct, or the genes combined in the vector construct are transferred together  
3 into selected recipients to obtain multiple gene-products or express multi-gene-  
4 depended characters.